



UNITED STATES PATENT AND TRADEMARK OFFICE

UNITED STATES DEPARTMENT OF COMMERCE United States Patent and Trademark Office Address: COMMISSIONER FOR PATENTS P.O. Box 1450 Alexandria, Virginia 22313-1450 www.uspto.gov

APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.	
10/053,753	01/22/2002	Lester F. Lau	05031.003.CNUS02	6127	
22930 7	590 07/22/2005		EXAM	EXAMINER	
HOWREY LI	- -		WOITACH,	JOSEPH T	
C/O IP DOCKETING DEPARTMENT 2941 FAIRVIEW PARK DR, SUITE 200			ART UNIT	PAPER NUMBER	
FALLS CHURCH, VA 22042-2924			1632		
			DATE MAIL ED: 07/22/2009	DATE MAILED: 07/22/2005	

Please find below and/or attached an Office communication concerning this application or proceeding.

Advisory Action						
Before the Filing of an Appeal Brief						

Application No.	Applicant(s)		
10/053,753	LAU, LESTER F.		
Examiner	Art Unit		
Joseph T. Woitach	1632		

3	LAGIIIIII	Airoille					
	Joseph T. Woitach	1632					
The MAILING DATE of this communication appe	ears on the cover sheet with the c	correspondence add	ress				
THE REPLY FILED <u>11 July 2005</u> FAILS TO PLACE THIS APF		-					
The reply was filed after a final rejection, but prior to or on the same day as filing a Notice of Appeal. To avoid abandonment of this application, applicant must timely file one of the following replies: (1) an amendment, affidavit, or other evidence, which places the application in condition for allowance; (2) a Notice of Appeal (with appeal fee) in compliance with 37 CFR 41.31; or (3) a Request for Continued Examination (RCE) in compliance with 37 CFR 1.114. The reply must be filed within one of the following time periods:							
The period for reply expires 3 months from the mailing date of the final rejection. The period for reply expires on: (1) the mailing date of this Advisory Action; or (2) the date set forth in the final rejection, whichever is later. In no							
event, however, will the statutory period for reply expire later than SIX MONTHS from the mailing date of the final rejection. Examiner Note: If box 1 is checked, check either box (a) or (b). ONLY CHECK BOX (b) WHEN THE FIRST REPLY WAS FILED WITHIN TWO MONTHS OF THE FINAL REJECTION. See MPEP 706.07(f).							
Extensions of time may be obtained under 37 CFR 1.136(a). The date on been filed is the date for purposes of determining the period of extension a CFR 1.17(a) is calculated from: (1) the expiration date of the shortened stabove, if checked. Any reply received by the Office later than three month earned patent term adjustment. See 37 CFR 1.704(b). NOTICE OF APPEAL	which the petition under 37 CFR 1.136(a and the corresponding amount of the fee. atutory period for reply originally set in the	The appropriate extension final Office action; or (2)	on fee under 37 as set forth in (b)				
2. ☐ The Notice of Appeal was filed on A brief in com	pliance with 37 CER 41 37 must be	a filed within two mon	the of the date				
of filing the Notice of Appeal (37 CFR 41.37(a)), or any e Since a Notice of Appeal has been filed, any reply must l	extension thereof (37 CFR 41.37(e)), to avoid dismissal (of the appeal.				
AMENDMENTS	Landa and the Albanda and Alba	8 - 20 - 41 - 4 - 1	•				
 The proposed amendment(s) filed after a final rejection, (a) They raise new issues that would require further contains. 	•		because				
(b) They raise the issue of new matter (see NOTE below		TE Delow),					
(c) They are not deemed to place the application in be appeal; and/or	•	educing or simplifying	the issues for				
(d) They present additional claims without canceling a NOTE: (See 37 CFR 1.116 and 41.33(a))		jected claims.					
4. The amendments are not in compliance with 37 CFR 1.	121. See attached Notice of Non-C	ompliant Amendment	(PTOL-324).				
5. $igsqcup$ Applicant's reply has overcome the following rejection(s							
 Newly proposed or amended claim(s) would be a the non-allowable claim(s). 	·	-					
7. For purposes of appeal, the proposed amendment(s): a) how the new or amended claims would be rejected is pro The status of the claim(s) is (or will be) as follows:		vill be entered and an	explanation of				
Claim(s) allowed:							
Claim(s) objected to:			•				
Claim(s) rejected: <u>65 and 67-77</u> .							
Claim(s) withdrawn from consideration:							
AFFIDAVIT OR OTHER EVIDENCE		.1.4:					
 The affidavit or other evidence filed after a final action, b because applicant failed to provide a showing of good ar and was not earlier presented. See 37 CFR 1.116(e). 	nut before or on the date of filing a find sufficient reasons why the affida	vit or other evidence	is necessary				
9. The affidavit or other evidence filed after the date of filing entered because the affidavit or other evidence failed to showing a good and sufficient reasons why it is necessar	overcome <u>all</u> rejections under appe	al and/or appellant fa	ils to provide a				
10. 🔲 The affidavit or other evidence is entered. An explanation							
REQUEST FOR RECONSIDERATION/OTHER							
11. The request for reconsideration has been considered but does NOT place the application in condition for allowance because: <u>See Continuation Sheet.</u>							
12. \square Note the attached Information Disclosure Statement(s).	(PTO/SB/08 or PTO-1449) Paper	No(s)	8				
13. Other:			7				
:	(< local.	hales -				
		Joe h					
		\sim	1150				

AU1632

÷

Continuation of 11. does NOT place the application in condition for allowance because:

Applicants argue that the specification provides guidance for making an antibody that specifically binds to human Cyr61 and can be distinguished from the cited art. Applicants arguments have been fully considered but not found persuasive. Upon review of the citation for support of Applicants' arguments, it is noted that the citation is not complete.

"These sequence differences are exploited to elicit antibodies specific to the human Cyr6l by using as an antigen a peptide having a sequence derived from one of the divergent regions in the human protein, although antibodies directed to a conserved region are also contemplated by the invention."

Clearly the specification provides support for using the divergent region(s) of Cyr61, however it clearly teaches that the conservered regions are a contemplated part of the invention as demonstrated by the complete sentence Applicants have cited. Examiner would not contest that differences between the mouse and human Cyr61 sequences exist, rather at issue is what weight the term "specific" should be given in the instantly claimed invention. Presently, the claim is being interpreted to encompass the fact that the antibody will bind Cry61 and not any other protein(s) non-specfically. Further, it should be noted that binding is usually empiracally tested and that the concentration of a given antibody solution and/or washing conditions would affect the "specific" binding of any antibody. Even if claims were to be interpreted to encompass antibodies that only bound the divergent sequence found between mouse and human, it is noted that the only teaching in the present specification is differences between specific mouse and human sequences, and fails to proivde any guidance to sequences from other species and human/. For the reasons above and of record, the rejection is maintained.